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Date of Filing

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Application number

200202366-1

Applicants

INSTITUTE OF MATERIALS RESEARCH

AND ENGINEERING

Title of Invention

METHOD

FOR

ELECTROLESS

PCT

WIPO

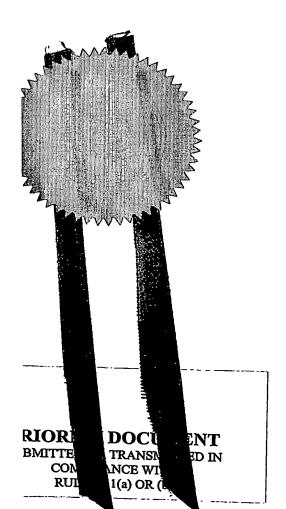
DEPOSITION OF A METAL LAYER ON

SELECTED PORTIONS OF A SUBSTRATE

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26 April 2003





PATENTS FORM 1

Patents Act (Cap. 221) Patents Rules Rule 19

INTELLECTUAL PROPERTY OFFICE OF SINGAPORE



REQUEST FOR THE GRANT OF A PATENT UNDER

			SECTION 25			
* denotes mandatory fiel	ds					
1. YOUR REFERENCE*		A1-1036 P	PK/rz			
2. TITLE OF INVENTION*	METHOD FO	OR ELECTRO PORTIONS (OLESS DEPOSITION OF OF A SUBSTRATE	A METAL LA	YER ON	
3. DETAILS OF APPI	LICANT(S)* (see	note 3)	Number of applic	cant(s)	1]
(A) Name	INSTITUTE	OF MATERI	ALS RESEARCH AND E	NGINEERING	}	
Address	3 RESEARCE SINGAPORE	H LINK E 117602				
State				Country	SG	
X For corpora	te applicant		For individual	applicant		' 1
State of incorporation			State of residency			
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(B) Name						
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For corporate applicant For individual applicant
State of incorporation State of residency
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For others (please specify in the box provided below)
(C) Name
Address
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For corporate applicant For individual applicant
State of incorporation State of residency
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For others (please specify in the box provided below)
Further applicants are to be indicated on continuation sheet 1
DECLARATION OF PRIORITY (see note 5) A. Country/country designated
File number Filing Date
B. Country/country designated
File number Filing Date
Further details are to be Indicated on continuation sheet 6
5. INVENTOR(S)* (see note 6)
A. The applicant(s) is/are the sole/joint inventor(s) Yes No X
Patents Form 1 Page 2 of 5

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B. A sta	tement on Patents Form 8 is/ will b	e-furnished Y	Yes X No			
6. CLAI	MING AN EARLIER FILING DATE	UNDER (see note	7)			
		ection 26(6)	section 47(4)			
Patent a	pplication number					
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Please (Note: C	mark with a cross in the relevant cl only one checkbox may be crossed	neckbox provided be .)	olow .			
Proceedings under rule 27(1)(a) DD MM YYYY						
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	Proceedings under rule 27(1)(b)					
	ON 14(4)(C) REQUIREMENTS (S		es No X			
	n has been displayed at an interna	tional exhibition. Y	es No A			
8. SECT	ION 114 REQUIREMENTS (see r	ote 9)				
The inve	ention relates to and/or used a mic	m-omanism denosit	ed for the numbers of disclosure	In accordance with section 114 with		
a depos	tory authority under the Budapest	Treaty.	ed for the purposes of disclosure	in accordance with section 114 with		
Yes	No X					
9. CHE	CKLIST*	•				
(A) The	application consists of the followi	ng number of sheets	3	•		
i.	Request	5	Sheets			
ii.	Description	10	Sheets			
III.	Claim(s)	3	Sheets			
iv.	Drawing(s)	7 .	Sheets			
v.	Abstract (Note: The figure of the drawing, if any, should accompany the abstract)	1	Sheets			
Total nu	mber of sheets	26	Sheets			
(B) The application as filed is accompanied by:						
	Priority document(s)	T	ranslation of priority document(s)			

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X Statement of inventorship & right to grant International exhibition certificate					
10. DETAILS OF AGENT (see notes 10, 11 and 12)					
Name					
Firm ARTHUR LOKE BERNARD RADA & LEE					
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12. NAME, SIGNATURE AND DECLARATION (WHERE APPROPRIATE) OF APPLICANT OR AGENT* (see note 12) (Note: Please cross the box below where appropriate.) I, the undersigned, do hereby declare that I have been duly authorised to act as representative, for the purposes of this					
application, on behalf of the applicant(s) named in paragraph 3 herein. DD MM YYYY 23/04/2002					
Name and Signature					

2 3 APR 2002 200202366-1



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- The relevant checkboxes as indicated in bold should be marked with a cross where applicable.
- Enter the name and address of each applicant in the spaces provided in paragraph 3. Where the applicant is an individual
 - · Names of individuals should be indicated in full and the surname or family name should be underlined.
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Where the applicant is a partnership

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- The checkbox for "For others" should be marked with a cross and the name and address of the partnership should be indicated in the box provided.
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- The declaration of priority in paragraph 4 should state the date of the previous filling, the country in which it was made, and indicate the file number, if available. Where the application relied upon in an International Application or a regional patent application e.g. European patent application, one of the countries designated in that application [being one falling under section 17 of the Patents Act] should be identified and the country should be entered in the space provided.
- Where the applicant or applicants is/are the sole inventor or the joint inventors, paragraph 5 should be completed by marking with a cross the 'YES' checkbox in the declaration (A) and the 'NO' checkbox in the alternative statement (B). Where this is not the case, the 'NO' checkbox in declaration (A) should be marked with a cross and a statement will be required to be filed on Patents Form 8.
- When an application is made by virtue of section 20(3), 26(6) or 47(4), the appropriate section should be identified in paragraph 6 and the number of the earlier application or any patent granted thereon identified. Applicants proceeding under section 26(6) should identify which provision in rule 27 they are proceeding under. If the applicants are proceeding under rule 27(1)(a), they should also indicate the date on which the earlier application was amended.
- Where the applicant wishes an earlier disclosure of the invention by him at an International Exhibition to be disregarded in accordance with section 14(4)(c), then the 'YES' checkbox at paragraph 7 should be marked with a cross. Otherwise, the 'NO' checkbox should be marked with a cross.
- Where in disclosing the invention the application refers to one or more micro-organisms deposited with a depository authority under the Budapest Treaty, then the 'YES' checkbox at paragraph 8 should be marked with a cross. Otherwise, the 'NO' checkbox should be marked with a cross. Attention is also drawn to the Fourth Schedule of the Patents Rules.
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METHOD FOR ELECTROLESS DEPOSITION OF A METAL LAYER ON SELECTED PORTIONS OF A SUBSTRATE

Technical Field

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The present invention relates to a method for electroless deposition of a metal layer on selected portions of a substrate. A preferred form of the invention relates to a method of depositing a desired metal layer, by electroless deposition, on one or more selected portions of an indium tin oxide (*ITO*) surface of a substrate. These selected portions are typically transparent conductive paths of ITO.

Background to the Invention

In this specification, where a document, act or item of knowledge is referred to or discussed, this reference or discussion is not an admission that the document, act or item of knowledge or any combination thereof, was at the priority date:

- (a) part of the common general knowledge; or
- (b) known to be relevant to any attempt to solve any problem with which this specification is concerned.
- Wet chemical metallisation of semiconductors has substantial industrial and commercial significance. This metallisation is generally effected by means of electroless plating on a suitable substrate. Such substrates typically include metals, ceramics (including glass) and polymers.
 - When, for example, it is desired to deposit nickel on such a substrate by electroless deposition, the substrate is typically immersed in a solution containing a reducing agent and nickel ions (Ni²⁺). The nickel ions are reduced by the reducing agent, usually in the presence of a suitable catalyst. It is usual practice to catalytically activate the surface of the substrate which is to be plated so as to confine deposition of the metal to the desired substrate surface.
- 30 A common catalyst used in this reaction is palladium.

However, it is often difficult to selectively deposit the desired metal layer on selected portions of the substrate surface. This is because, when a substrate surface is activated, the metal will tend to deposit on all of the activated surface. Therefore, it may then be necessary to deactivate portions of the surface where the metal is not to be deposited.

For example, US 4824693 discloses a method of depositing a solderable metal layer, by an electroless process, on transparent conductive paths of ITO on certain substrates (including glass substrates). The invention of US 4824693 involves activating the surface of the conductive paths by immersing the entire substrate surface in a bath of palladium chloride and tin chloride then deactivating the areas not covered by the conductive paths (by immersing the substrate in hydrofluoric acid). The process then includes deposition of the metal layer by an electroless method. The metal will deposit on the remaining activated portions of the surface. As can be seen, in the method of US 4824693, the activation of the surface is not selective and, therefore, portions of the ITO surface must then be deactivated prior to the metal deposition.

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European Patent No. 0 518 422 A2 discloses a method for selectively activating an ITO surface of a glass substrate. This patent uses polymer-stabilised colloidal particles that selectively adsorb to the ITO surface and do not adsorb to the glass substrate. However, this patent only describes how to activate all of the ITO surface of the substrate and it does not contain any indication as to how to selectively activate desired portions of the ITO surface.

The present invention is directed towards an improved method for selectively activating a substrate so as to facilitate metallic deposition thereon.

This invention is further directed towards an improved method of selectively activating and plating desired portions of a substrate, in particular, an ITO surface of a substrate.

Summary of the Invention

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According to a first aspect of this invention, there is provided a method for the electroless deposition of a desired metal layer on one or more selected portions of a substance surface, wherein the method includes the steps of:

- applying a masking layer onto the surface of the substrate, said masking layer adapted to have one or more apertures formed therein so as to expose the one or more selected portions of the surface;
 - exposing the one or more selected portions of the surface to a colloidal suspension of catalytic particles adapted to adsorb to the substrate surface and to enhance deposition of the desired metal layer thereon; and
 - exposing the one or more selected portions of the surface to an ionic solution containing ions of the desired metal to enable formation of the metal layer.

It is preferred that the one or more apertures are formed in the masking layer after applying the layer to the substrate surface.

It is further preferred that the substrate surface has a film of ITO formed thereon. This film may cover all of, or a portion of, the substrate surface. Alternatively, the film may be applied in patches to the substrate surface.

Preferably, at least some of the one or more apertures of the masking layer lie over one or more portions of the ITO film.

It is preferred that the colloidal suspension comprises particles of catalytic metal which are adapted to adsorb to the ITO film but not to the substrate material. One particularly preferred catalytic metal is palladium. A preferred substrate material is glass.

The suspension of catalytic metal particles is generally polymer-stabilised, so as to inhibit the metal particles from agglomerating and precipitating from the solution. This polymer stabilisation is preferably achieved using polyvinyl alcohol (PVA) or poly(vinylpyrrolidone) (PVP). The catalytic metal particles may also be stabilised with a solution containing tin ions or a combination of tin ions and the polymers mentioned above.

The masking layer may be formed of a photosensitive material. Suitable such masking layers are dry film resists which may be selected from Ashahi Chemical's SunfortTM resists. Alternatively, the dry film resist may be selected from DuPont's RistonTM resists. Alternatively, the masking layer may be formed of a polymeric material to which no substantial adherence of the catalytic particles occurs. Similarly, the catalytic particles may be selected on the basis that they do not (to any significant extent) adhere to the polymeric material. The polymeric material is typically selected from the group consisting of suitable polycarbonates, fluorinated polymers, cellophane, polyimide and acrylate-based polymers. The masking layer is preferably applied by coating the substrate surface with a thin layer of a photoresistive polymer. This coating can be achieved by spin coating a liquid resist on the substrate surface and subsequently curing and/or drying the liquid resist. Alternatively, the masking layer may be applied by spraying an aerosol of a suitable polymeric material (such as those described above) onto the substrate surface.

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Another means of applying the masking layer is to laminate a film of the resist onto the substrate surface. Particularly suitable films for this purpose are the dry film resists mentioned above. This lamination typically involves placing the film on top of the substrate and then feeding them between two rollers that apply pressure and heat causing the film and substrate to adhere to each other.

The one or more apertures in the masking layer may be formed either before of after the masking layer has been applied to the substrate. However, it is generally preferable to form the apertures after applying the masking layer to the substrate.

Typically, the apertures are formed using UV lithography (in the case of photosensitive materials), a laser or screening means. Preferably, a photosensitive material is used for the patterning. This UV lithography typically involves exposing the photoresist to UV light selectively, such as through a UV mask. The UV mask is usually a patterned thin layer of metal on glass.

30 Some resists are 'positive' meaning that the regions of such resists which are exposed to UV light are broken down and become soluble in the developing agent. Other resists are 'negative', meaning that the regions of such resists

which are exposed to UV light harden and become insoluble to the developing agent.

After exposure to UV light, the layered substrate is immersed in a suitable developing agent adapted to selectively remove the exposed regions of the resist (in the case of 'positive' resists) or the unexposed regions of the resist (in the case of 'negative' resists). This results in the desired patterns (of apertures) being formed in the masking layer.

It is preferred that, prior to performing the step of exposing the selected portions of the substrate to the colloidal solution, the layered substrate is cleaned to remove any residues of polymeric or organic material. This cleaning may be effected by plasma cleaning or by UV ozone cleaning techniques.

The step of exposing the one or more selected portions of the substrate to the colloidal suspension is generally effected by dipping the substrate, containing the masking layer, into a bath of the colloidal suspension. The substrate may be removed from the suspension after a few seconds, or longer if necessary. After exposing the selected portions to the colloidal suspension, the method preferably includes the further step of rinsing the selected portions, which is preferably done with deionized water (DI water). After the rinsing step, the selected portions should be dried. Drying can be effected by placing the layered substrate in an oven to remove substantially all of the water from the selected portions, and/or by blowing the substrate with a stream of gas, such as nitrogen or air.

The step of exposing the one or more selected portions to the ionic solution is generally effected by dipping the substrate containing the masking layer into a bath of the ionic solution.

It is preferred that, after the metal layer has been formed, the masking layer is removed. However, it is also possible for the masking layer to be removed after exposing the selected portions to the colloidal suspension but prior to exposing the one or more selected portions to the ionic solution. Removal of the masking layer is typically facilitated by using a strong base (such as

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The desired metal for deposition on the substrate is typically copper, nickel, chromium, molybdenum, tantalum or any alloys of these. Copper or nickel are particularly preferred as the desired metal.

In a particularly preferred embodiment of this invention, the method is effected by performing the following steps:

- applying a masking layer on to the surface of the substrate;
- forming the desired apertures in the masking layer;

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- dipping the substrate into the colloidal suspension of catalytic particles;
 - rinsing the substrate with clean water immediately after dipping it in the colloidal suspension;
 - drying the substrate (usually with flowing air or nitrogen) and then further drying it in an oven;
- immersing the substrate in an electroless metallisation bath until the desired thickness of the desired metal is deposited on the one or more selected portions of the surface of the substrate; and
 - stripping the masking layer from the substrate.

Additional steps may be included to further facilitate the method. For instance, prior to applying the masking layer, an ITO film (or portions of such a film) may be applied to the substrate surfaces.

The colloidal suspension referred to above contains catalytic particles, which are typically catalytic metals such as palladium. Preferably the colloidal solution is prepared as follows:

- dissolving 100mg (although it could be anywhere from about 25mg to about 500mg) of PVP (weight averaged molecular weight = 50,000 although it could be anywhere between about 10,000 to about 500,000) in DI water;
 - dissolving 150mg of PdCl₂ in 5.25ml of HCl (37% aqueous solution);
- mixing the PVP and the PdCl₂ solutions together;

- slowly adding 10-35ml of hypophosphorous acid (H₃O₂P) (50% aquoeus solution) to the solution (or an appropriate quantity of another suitable reducing agent);
- adding DI water until the total volume of the solution is about 1 litre.
- The PVP stabilised palladium colloidal solution (the "Pd/PVP sol") is very stable and can be kept for months under normal conditions without noticeable change. In preparing the substrate for the metal deposition, a clean substrate, having an ITO film or ITO pattern on its surface, is taken and coated with a masking layer (or "resist") of the type described above.
- The resist needs to be patterned so that the patterns correspond with the selected portions of the substrate on to which the desired metal layer is to be deposited. Preferably a photoresistive material is used for the patterning. Patterning of the photoresist on the substrate surface generally follows procedures known in the art. These include exposure of selected regions of the photoresist to UV light. Thereafter the resist is immersed in a developing agent to selectively dissolve and remove the resist. The exposed areas on the substrate should be cleaned after patterning of the resist so as to remove any residues of polymeric or organic material left on the surface of the substrate.
- The clean patterned substrate is then dipped into the Pd/PVP sol. Usually the substrate only needs to be dipped into the solution for a few seconds. Upon removal of the substrate from the solution, it should be immediately rinsed thoroughly with DI water. The substrate may then be dried with nitrogen or air and may then be placed in an oven to remove any remaining moisture.

The substrate is then placed in an electroless plating bath to deposit the desired thickness of metal on the exposed selected portions of the substrate.

Generally, the particles in the Pd/PVP sol do not adhere to the resist and, therefore, no plating generally occurs on the resist. The resist may then be removed from the substrate. Alternately, the resist may be removed from the substrate prior to the metal deposition step (but after the step of activating the substrate with the Pd/PVP sol). In certain cases, however, it may be desirable to leave the resist on the substrate.

Drawings

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Figure 1 is a flow chart of a method according to a preferred embodiment of this invention.

Figure 2 is a schematic perspective view of a substrate according to a preferred embodiment of this invention.

Figure 3 is a schematic perspective view of a substrate with masking layer (resist) applied thereto, according to a preferred aspect of this invention.

Figure 4A is a schematic perspective view of the substrate and masking layer of Figure 3 in which the masking layer is patterned to expose selected portions of the substrate, according to a preferred aspect of this invention.

Figure 4B is a schematic plan view of the substrate and masking layer of Figure 4A.

Figure 5 is a side cross-sectional view of a bath of colloidal solution into which the substrate and masking layer of Figures 4A and 4B is being dipped.

Figure 6 is a side cross-sectional view of a bath, containing metal ions of the metal to be deposited on the substrate, into which the substrate and masking layer is being dipped.

Figure 7 is a schematic side view of the substrate of Figures 4A and 4B with partially removed masking layer, following the metal deposition steps according to a preferred aspect of this invention.

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Figure 8 is a schematic perspective view of the substrate (with masking layer removed) after the metal deposition, according to a preferred embodiment of this invention.

Figure 9 is an optical microscope image of a partial plan view of the substrate and masking layer following metal deposition.

Figure 10 is an enlarged view of a portion of the substrate and masking layer of Figure 9.

25 Figure 11 is a further enlarged view of a portion of the substrate and masking layer of Figure 10.

Figure 12 is an optical microscope image of a partial plan view of the substrate, showing metal deposited on selected portions of the substrate (with masking layer removed).

Figure 13 is an enlarged view of a portion of the substrate of Figure 12.

Figure 14 is a further enlarged view of a portion of the substrate of Figure 12.

Brief Description of the Drawings

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As shown in Figures 2 and 3, the substrate 1 is a substantially rectangular support on an upper surface of which is located a film of ITO 2. A masking layer (resist) 3 is placed over the upper surface of the substrate.

Figure 3 shows the masking layer 3 as covering the entire upper surface of the substrate 1. This may be preferred, for example, in situations where the seeding particles (and therefore the deposited metal) may otherwise adhere to the substrate. However, where the seeding particles do not (at least, to a significant degree) adsorb to the substrate, the masking layer may be dimensioned so as only to cover the ITO film. As shown in Figures 4A and 4B, the masking layer 3 is patterned with shaped holes 4. Shaped holes 4 are dimensioned so as to correspond with the selected portions of the substrate on to which metal is to be deposited.

Figure 5 shows the layered substrate 1 being dipped into a bath 5 containing a colloidal suspension 6 of catalytic metal particles, namely palladium.

Figure 6 shows the layered substrate 1 being dipped into a bath 7 of an ionic solution 8 containing ions of the desired metal (eg. nickel).

Figure 7 shows the layered substrate 1, with the masking layer 3 partially removed from the substrate 1, following the step of metal deposition and showing a metal portion 9 deposited on the substrate 1.

Figure 8 shows the substrate 1 having metal portions 9 deposited on the selected portions of the substrate 1.

Figures 9 and 10 show a portion of the upper surface of the substrate 1 following metal deposition but prior to removal of the masking layer 3. The masking layer 3 contains a multitude of shaped holes (most of which are circular 4a and one of which is linear 4b). In the embodiment shown in Figures 9 and 10, a clear dividing line 10 can be seen — above which no metal has been deposited and below which metal has been deposited. Below the dividing line 10, metal can be seen to have been deposited in the shaped holes of the masking layer 3. This deposition occurred because of the presence of the seeded ITO film on the substrate.

Figure 10 is an enlarged version of a portion of Figure 9 showing the portion of the substrate surface (with masking layer) near the dividing line 10. As can be seen, in the shaped holes 4a, 4b which are above the dividing line 10, being those which overlay portions of the substrate on which the ITO film had not

been deposited, the metal did not adhere. This is also shown in Figures 11, 12, 13 and 14 (in all of which the masking layer 3 has been removed). Figure 11 shows a single circular shaped hole 4a which contains a metallised section 20 and a non-metallised section separated by the dividing line 10.

The invention described above provides a selective method of seeding and plating only certain portions of a substrate having an ITO film. This enables simple, accurate patterning (eg. on the microscale) of the substrate.

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Modifications and improvements to the invention will be readily apparent to those skilled in the art. Such modifications and improvements are intended to be within the scope of this invention.

Claims

The claims defining this invention are as follows:

- 1. A method for the electroless deposition of a desired metal layer on one or more selected portions of a substrate surface, wherein the method includes the steps of:
 - applying a masking layer onto the surface, said masking layer adapted to have one or more apertures formed therein so as to expose the one or more selected portions of the surface;
 - exposing the one or more selected portions of the surface to a colloidal suspension of catalytic particles adapted to adsorb to the substrate surface and to enhance deposition of the desired metal layer thereon; and
 - exposing the one or more selected portions of the surface to an ionic solution containing ions of the desired metal to enable formation of the metal layer.
- 2. A method according to claim 1, wherein the one or more apertures are formed in the masking layer after applying the layer to the substrate surface.
- 3. A method according to claim 1 or claim 2, wherein the substrate has a film of indium tin oxide (ITO) formed thereon.
 - 4. A method according to claim 3 wherein at least some of the one or more apertures of the masking layer lie over one or more portions of the ITO film.
- 5. A method according to any one of claims 1 to 4, wherein the colloidal suspension includes particles of catalytic metal.
 - 6. A method according to claim 5, wherein, when the substrate surface includes a film of ITO formed thereon, the catalytic metal and the material of the substrate are selected so that no substantial adsorption of the catalytic metal occurs on the substrate material.
- 30 7. A method according to claim 5 or claim 6, wherein the catalytic metal is palladium.
 - 8. A method according to any one of claims 5 to 7, wherein the catalytic metal particles are polymer-stabilised.

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- A method according to claim 8, wherein the catalytic metal particles are stabilised with polyvinyl alcohol, poly(vinylpyrrolidoine) or a combination of these.
- 10. A method according to any one of claims 5 to 7, wherein the catalytic metal particles are stabilised with a solution containing tin ions.
 - 11. A method according to any one of claims 6 to 10, wherein the substrate material is glass.
 - 12. A method according to any one of claims 1 to 11, wherein the masking layer is formed of a polymeric material to which no substantial adherence of the catalytic particles occurs.

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- 13. A method according to claim 12, wherein the polymeric material is selected from the group consisting of suitable polycarbonates, fluorinated polymers, cellophane, polyimide and acrylate-based polymers.
- 15 14. A method according to claim 12 or claim 13, wherein the polymeric material is a photoresist.
 - 15. A method according to any one of claims 1 to 14, wherein the masking layer is formed of a dry film resist.
- 16. A method according to claim 15, wherein the dry film resist is selected from the group consisting of Asahi Chemical's SunfortTM resists and DuPont's RistonTM resists.
 - 17. A method according to any one of claims 12 to 16, wherein the one or more apertures in the masking layer are formed using UV lithography, a laser or screening means.
- 25 18. A method according to any one of claims 1 to 17 wherein, prior to the step of exposing the selected portions of the substrate to the colloidal solution, the layered substrate is cleaned to remove any residues of polymeric or organic material.
- 19. A method according to claim 18, wherein the cleaning is effected by plasma cleaning or UV ozone cleaning techniques.
 - 20. A method according to any one of claims 1 to 19, wherein the step of exposing the one or more selected portions of the substrate to the colloidal solution is effected by dipping the substrate containing the masking layer into a bath of the colloidal solution.

- 21. A method according to any one of claims 1 to 20, wherein, after the step of exposing the one or more selected portions of the substrate to the colloidal solution, the selected portions are rinsed with de-ionised water.
- 22. A method according to claim 21 wherein, after the rinsing step, the selected portions are dried to remove substantially all of the water from the selected portions.

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- 23. A method according to claim 22, wherein the drying step includes placing the layered substrate in an oven.
- 24. A method according to claim 22, wherein the drying step includesblowing a stream of gas over the layered substrate.
 - 25. A method according to any one of claims 1 to 25, wherein the drying step includes both placing the layered substrate in an oven and blowing it with a stream of gas.
- 26. A method according to claim 24, wherein the step of exposing the one or more selected portions to the ionic solution is effected by dipping the substrate containing the masking layer into a bath of the ionic solution.
 - 27. A method according to any one of claims 1 to 26 wherein, after formation of the metal layer, the masking layer is removed.
- 28. A method according to claim 27 wherein a strongly basic solution is used to facilitate removal of the masking layer.
 - 29. A method according to any one of claims 1 to 25, wherein the masking layer is removed prior to the step of exposing the one or more selected portions to the ionic solution.
- 30. A method according to any one of claims 1 to 29, wherein the desired metal is selected from the group consisting of copper, nickel, chromium, molybdenum, tantalum and any alloy of these metals.
 - 31. A method according to claim 30, wherein the desired metal is selected from copper and nickel.
- 32. A method for the electroless deposition of a desired metal layer on one or more selected portions of a substrate surface, substantially as herein before described with reference to any one or more of the drawings.
 - 33. A product made according to the method of any one or more of claims 1 to 32.

Abstract

Method for Electroless Deposition of a Metal Layer on Selected Portions of a Substrate

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The present invention relates to a method for electroless deposition of a metal layer on selected portions of a substrate. A preferred form of the invention relates to a method of depositing a desired metal layer, by electroless deposition, on one or more selected portions of an indium tin oxide (ITO) surface of a substrate. These selected portions are typically transparent conductive paths of ITO. The method includes a number of steps. The first step involves applying a masking layer onto the surface, the masking layer has one or more apertures formed therein so as to expose the one or more selected portions of the surface. The next step involves exposing the one or more selected portions of the surface to a colloidal suspension of catalytic particles adapted to adsorb to the substrate surface and to enhance deposition of the desired metal layer thereon. The next step involves exposing the one or more selected portions of the surface to an ionic solution containing ions of the desired metal to enable formation of the metal layer.

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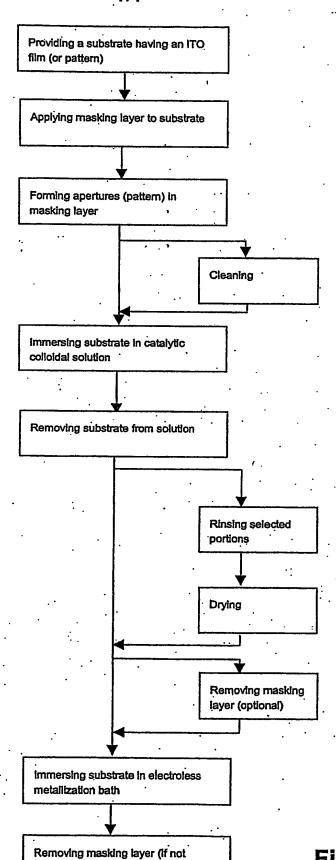
[Figure 1]







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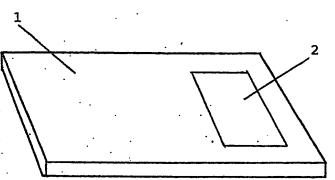
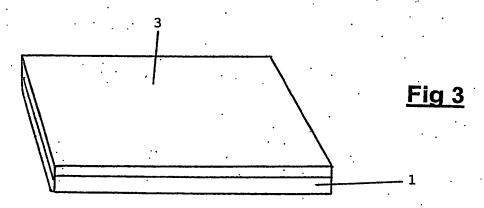
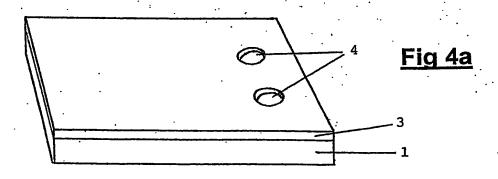


Fig 2





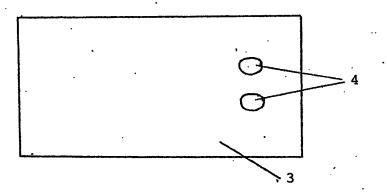
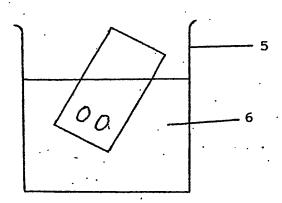


Fig 4b



<u>Fig 5</u>

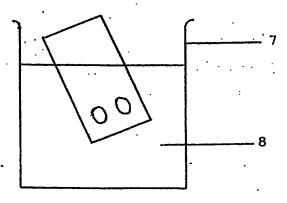
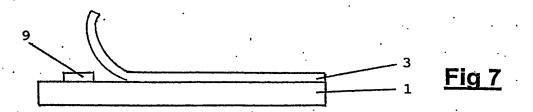
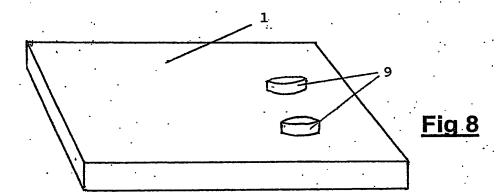
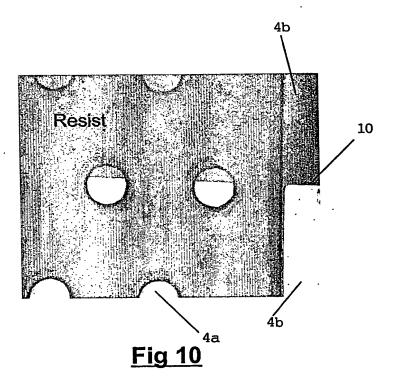
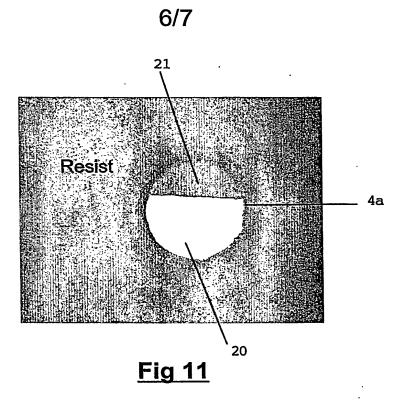


Fig 6









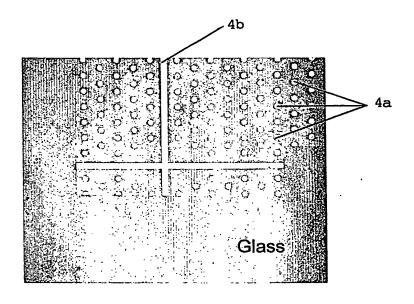


Fig 12

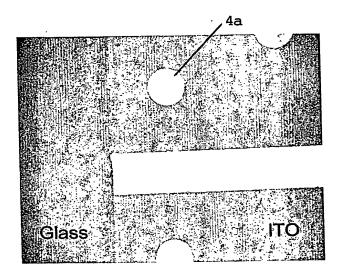


Fig 13

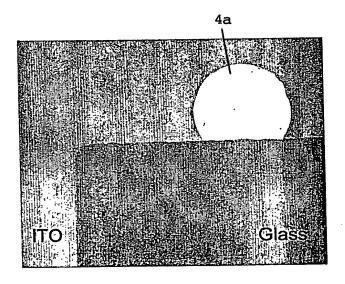


Fig 14

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